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Could the identification of subclinical atherosclerosis offer an alternative to the mass drug treatment of hypercholesterolemia?

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*Short communication*

Identification of the homozygous missense mutation in the lecithin:cholesterol-acyltransferase (LCAT) gene, causing LCAT familial deficiency in two French patients

I. Dorval, P. Jezequel, C. Dubourg, B. Chauvel, P. Le Pogamp, J.-Y. Le Gall (France)

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